

ENCODE data at the ENCODE portal

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ABSTRACT

The Encyclopedia of DNA Elements (ENCODE) Project is in its third phase of creating a comprehensive catalog of functional elements in the human genome. This phase of the project includes an expansion of assays that measure diverse RNA populations, identify proteins that interact with RNA and DNA, probe regions of DNA hypersensitivity, and measure levels of DNA methylation in a wide range of cell and tissue types to identify putative regulatory elements. To date, results for almost 5000 experiments have been released for use by the scientific community. These data are available for searching, visualization and download at the new ENCODE Portal (www.encodeproject.org). The revamped ENCODE Portal provides new ways to browse and search the ENCODE data based on the metadata that describe the assays as well as summaries of the assays that focus on data provenance. In addition, it is a flexible platform that allows integration of genomic data from multiple projects. The portal experience was designed to improve access to ENCODE data by relying on metadata that allow reusability and reproducibility of the experiments.

INTRODUCTION

The Encyclopedia of DNA Elements (ENCODE) Project began as a Pilot Project on 1% of the human genome (1,2). In 2007, the effort was scaled to whole-genome assays followed by expansion to performing similar assays in mouse (3–5). The ENCODE Project continues to create a comprehensive catalog of gene elements and functional elements in the human and mouse genomes by measuring RNA expression levels, identifying proteins that interact with RNA and DNA (such as modified histones, transcription factors and RNA-binding proteins), measuring

the levels of DNA methylation and identifying regions of DNA hypersensitivity. These data generated by ENCODE Consortium members are submitted to the ENCODE Data Coordination Center (DCC) whose primary task is to curate, process and validate the data in preparation for release to the scientific community. To promote the sharing of data, the ENCODE consortium has updated its release policy (<https://www.encodeproject.org/about/data-use-policy/>). These data are made available at the ENCODE Portal (www.encodeproject.org), created by the DCC, and are also distributed through the UCSC genome browser, GEO and Ensembl and used extensively as a community resource (6–11).

To increase the reproducibility, reusability and interoperability of the ENCODE data, the ENCODE DCC, in collaboration with the ENCODE consortium, has defined a new metadata standard. The experimental assays and computational methods used to generate the ENCODE data are represented by a structured data model to capture metadata that allows for maximal understanding and interpretation of these results (Hong *et al.*, submitted). Integral to the metadata, persistent identifiers are assigned for experiments, files, antibody lots and biosamples to allow specific identification of the data. In addition, controlled vocabularies and ontologies are used to increase the interoperability of the ENCODE data with other projects (12).

Specifically, the experiment metadata displayed on the portal utilize the Uber Anatomy Ontology (UBERON, uberon.org) for tissue biosamples; Cell Ontology (CL, cellontology.org) for primary cells; Experimental Factor Ontology (EFO, www.ebi.ac.uk/efo/) for immortalized cell lines; Ontology for Biomedical Investigations (OBI, obiontology.org) for experimental assays; Chemical Entities of Biological Interest (ChEBI, www.ebi.ac.uk/chebi/) for chemical treatments; and Sequence Ontology (SO, sequenceontology.org) for the nucleic acids of interest.

As discussions about the issues of reproducibility and reusability of genomic data are becoming more significant (13,14), the new ENCODE Portal serves to communicate

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The image shows the ENCODE Portal home page with several annotations:

- (a) Points to the 'Data' menu item in the top navigation bar.
- (b) Points to the 'Methods' section, which lists 'Data standards', 'Software tools', 'Pipelines', and 'Experiment guidelines'.
- (c) Points to the 'About' section, which includes 'Project overview', 'News', 'Publications', 'Release policy', and 'Data access'.
- (d) Points to the 'Help' section, which lists 'Getting started', 'REST API', 'File formats', 'Ontologies', 'Tutorials', and 'Contact'.
- (e) Points to the search box in the top right corner.
- (f) Points to the 'Quick Start' section, which provides instructions on how to find and download data.
- (g) Points to the 'News' section, which displays updates and data releases.

Figure 1. Overview of the ENCODE Portal ‘dashboard-style’ home page. (a) The Data section provides links into search pages that allow filtering of the ENCODE data. (b) The Methods section contains standard documents, software and analysis pipeline information. (c) The About ENCODE section provides context to the ENCODE project, the site and access to relevant publications. (d) The Help section lists tutorials and guide pages. (e) The free-text search box for querying any metadata or text item. (f) The Quick Start guide has popular links to browsing and download information. (g) The News feed tracks updates and data released. Items (a) to (e) are part of a persistent menu bar that appears on every page to provide stable navigation throughout the site.

to the scientific community the experimental standards and guidelines defined by the entire Consortium, as well as the details of how the assays were performed and the computational methods used to analyze the data.

NEW ENCODE DATA PORTAL

The new ENCODE data portal, released in August 2014, is the canonical, central source for ENCODE raw data, analysis data, methods, standards and experimental metadata. The user experience begins with a dashboard-style home page (Figure 1), that includes a news feed, menus and a quick start guide providing instruction on the various ways to access the data via browsing, searching, visualizing and downloading. A persistent standard menu at the top of every page provides stable navigation throughout the site. Within the menu are the categories of ‘Data’, ‘Methods’, ‘About ENCODE’ and ‘Help’. The ‘Data’ section provides initial search access into the data as well as a link to the new Release Policy that describes the terms of data use. The primary access to the ENCODE data is through the ‘Assays’ search page. This page allows one to browse replicated experimental sets, by faceted search selection on the left to filter experiments by features such as assay type, biosample type, organism or treatment. The biosample page provides similar browsing access to a listing of all individual biosam-

ples used in the experiments. The antibody page provides access to the antibodies screened or used in the ENCODE immunoprecipitation experiments along with the required antibody characterizations generated by the project (15). The annotation section links publications and analysis files generated by the consortium towards the goal of creating an encyclopedia of DNA elements. This section also contains links to multiple browsers to view annotations created from collations of information from the many ENCODE experiments.

The ‘Methods’ section includes information on the data standards, software, analysis pipelines and experimental guidelines defined by the consortium. The ENCODE consortium develops internal standards on the quality and procedures for both the experimental methods and the computational analysis. Current standards and guidelines are available from the portal as well as an archive containing previous versions. Additionally, a large variety of software is used in the analysis of these datasets. To improve transparency, the ENCODE Portal is hosting version information and links to source code for the software used in ENCODE project analysis. The goal is to provide sufficient information to allow anyone in the community to easily repeat the analysis if so desired. The ENCODE consortium is also developing uniform analysis pipelines that map raw sequencing data for various assays, generate vi-

The figure illustrates the ENCODE portal's search and filtering capabilities. It shows three stages of a search process:

- Top Panel:** A search for 'heart' yields 105 experiments. The search bar contains 'heart' and a search icon. A 'keyword search' label with a red arrow points to the search bar.
- Middle Panel:** Filtering by 'RNA-seq' and 'Homo sapiens' results in 8 experiments. The 'Assay' facet shows 'RNA-seq' selected with 8 experiments. The 'Organism' facet shows 'Homo sapiens' selected with 8 experiments.
- Bottom Panel:** A list of 8 experiments is displayed, including:
 - RNA-seq of heart (*Homo sapiens*, fetal)
 - RNA-seq of heart (*Homo sapiens*, adult 34 year)
 - RNA-seq of heart (*Homo sapiens*, fetal)
 - RNA-seq of heart (*Homo sapiens*, adult 30 year)
 - RNA-seq of cardiac atrium fibroblast (*Homo sapiens*, child 2 year)
 - RNA-seq of cardiac ventricle fibroblast (*Homo sapiens*)
 - RNA-seq of endothelial cell of coronary artery (*Homo sapiens*, adult)
 - RNA-seq of smooth muscle cell of the coronary artery (*Homo sapiens*, adult)

Faceted browsing is highlighted on the left side of the middle panel, showing various filters like Assay, Experiment status, Genome assembly, Organism, Biosample type, Organ, Life stage, Pipeline, and Available data.

Figure 2. Using search and filtering to select relevant experiments. The organization of key experiment metadata into facets allows the user to quickly filter from nearly 5000 to only eight experiments with a few simple terms. Entering 'heart' in the search box reduced the list to 105 experimental datasets. Filtering further on the terms 'RNA-seq' and '*Homo sapiens*' using the facets limited the list even further down to 8 experiments.

visualizable signal files from those mappings and produce a quantification or annotation result. Summaries of these pipelines, along with general specifications, are linked from the 'Pipelines' page. The 'About ENCODE' section gives context to the wealth of experiments found on this site, providing a project overview, access to all news items and information about data access at other sites. This section also has the 'Publications' page which provides guided entry into browsing a curated collection of publications that either use ENCODE data or are references for the methods used in creating the ENCODE data. Finally, the 'Help' menu has a variety of help documents and tutorials to facilitate differing user experiences. Included in this section are instructions on how to browse the data in the UI, using the REST API programmatic download.

DATA ACCESS FEATURES

Access to the ENCODE data from the new portal is driven by high-quality metadata that provide data provenance and transparency by describing how the assays were performed and how the data were analyzed. In order to import the data from previous phases of the ENCODE Project, a massive

data review was undertaken to match files with controls, to include any information that was found in the documentation, to work with laboratory groups to bring their older metadata up to the newer standards (including capturing protocols, sequencing platforms used, replicate structure, and shared biosamples when it could be known), to assign specific ontology terms for the biosamples and capture library preparation details. With this depth and breadth of metadata, the new portal is able to provide a more advanced level of searching, a more comprehensive view of the assays performed and visualization of the ENCODE data using the selected subset of the data.

Browsing and searching

ENCODE data can be accessed both by browsing via faceted search and by searching via keywords. The faceted search page, accessible via the 'Assay' link underneath the 'Data' menu item in the toolbar, provides a simplified interface that allows the filtration of thousands of assays to the ones of interest in a few clicks. Using a few simple terms like heart, RNA-seq and *Homo sapiens* can reduce

such as the target of the antibody in a ChIP-seq assay, a summary of the biosample and biological replicate number. This feature allows the user to easily integrate ENCODE data with their own custom tracks or the many reference tracks available at the Genome Browser. As well as being on the individual experiment page, this feature is available on the Assay search page. Once the user has filtered the list to a reasonable subset (currently <500), the 'Visualize' button will appear and a track hub including all appropriate files from every experiment on the list is assembled.

FUTURE WORK

Here, we have described the new features of the ENCODE Portal: improved access to the experiment standards and methods developed by the ENCODE Consortium, metadata-driven searches to find relevant datasets and visualization of only relevant datasets. The ENCODE Portal will be continuously updated with new experimental data, processed data and analysis metadata along with enhanced searching and visualization features as the project continues. Specifically, we are planning on introducing the concept of series organization of experiments into our Assay page. With this feature, we will be able to include a collection of assays in a series of time points or differentiation stages in the faceted search along with replicated experimental sets. And to further improve searching, we are implementing ways to search the data files themselves for genomic regions, in order to direct the scientific community to relevant assays based on whether a binding site is present near a gene of interest or a gene is expressed in a specific tissue.

In addition to providing improved access to the ENCODE data, the ENCODE Portal can serve as a gateway for other genomic projects. We are currently integrating metadata and data from other epigenetics projects into the site. Specifically we will be hosting the data of the recently funded Genomics of Gene Regulation project (<http://www.genome.gov/27561317>), the published datasets of the modENCODE integration papers (18–20) and Roadmap Epigenomics Mapping Consortium (<http://www.roadmapepigenomics.org/>) (21). This will allow searching across projects for related assays. The basic framework for the site was built to allow complex relations of experimental metadata and it is made available for other projects. It is currently being adopted by the Clinical Genome Resource (ClinGen) (22).

We have worked with GEO at NCBI to map our biosample schema to their Biosample standard (www.ncbi.nlm.nih.gov/biosample) (11,23). Currently, we are working with a method to not simply import these data into their database, but also to periodically synchronize the two databases to provide access to the most up-to-date data produced by the ENCODE consortium to the larger scientific community.

CONTACT INFORMATION

General questions about the ENCODE data portal should be directed to the encode-help@lists.stanford.edu mailing list. For any specific questions about laboratory details or methods, the producing laboratory information

is available on each experimental page and in the metadata for each file. All of the source code used for the portal website is open source and available through our repository hosted at GitHub (www.github.com/ENCODE-DCC/). You can sign up for an ENCODE announcement mailing list at <https://mailman.stanford.edu/mailman/listinfo/encode-announce>. Data and portal updates are available on our news page at www.encodeproject.org/news/. You can also follow us on twitter at @encodedcc.

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